



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁶ : A61L 2/20, B01J 19/08	A1	(11) International Publication Number: WO 99/32162 (43) International Publication Date: 1 July 1999 (01.07.99)
(21) International Application Number: PCT/US98/26232 (22) International Filing Date: 22 December 1998 (22.12.98) (30) Priority Data: 60/068,668 23 December 1997 (23.12.97) US (71)(72) Applicants and Inventors: CARMEN, Gary, B. [US/US]; 5815 Lone Horse Drive, Reno, NV 89502 (US). WIRTZ, Stephen, K. [US/US]; 2230 Fignon Ranch Road, Sparks, NV 89434 (US). (74) Agents: DONNELLY, Laura, A. et al.; McDermott, Will & Emery, 600 13th Street, N.W., Washington, DC 20005-3096 (US).		(81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i>
(54) Title: DYNAMIC O ₂ BIOLOGICAL BURDEN REDUCTION (57) Abstract A method and apparatus for significantly reducing the biological load on consumer products such as food products, botanicals, cosmetic ingredients and medical products is disclosed. The method involves applying a continuous stream of O ₂ gas to a material in a sealed biological burden reduction chamber (1). The continuous stream of O ₂ gas is prepared in an O ₂ generation cell (6), which contains a means for generating the O ₂ gas at pressure less than 20 lbs/in ² using, for example, one or more of the following: corona discharge, high frequency electrical discharge, ultraviolet light, x-ray, radioactive isotope and electric beam. The apparatus contains: (a) a biological burden reduction chamber (1); (b) a vacuum pump (50) coupled to the biological burden reduction chamber (1); (c) an O ₂ generating cell (6), wherein the O ₂ generation cell (6) contains an O ₂ generator capable of generating O ₂ at a pressure less than 20 lbs/in ² ; (d) a first control valve coupled to the biological burden reduction chamber (1) and the O ₂ generation cell, wherein the first control valve is capable of permitting O ₂ to be drawn from the O ₂ generation cell (6) into the biological burden reduction chamber (1); and (e) a second control valve coupled to the biological burden reduction chamber (1), wherein the second control valve is capable of withdrawing O ₂ contained within the biological burden reduction chamber (1).		

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece	ML	Mali	TR	Turkey
BG	Bulgaria	HU	Hungary	MN	Mongolia	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MR	Mauritania	UA	Ukraine
BR	Brazil	IL	Israel	MW	Malawi	UG	Uganda
BY	Belarus	IS	Iceland	MX	Mexico	US	United States of America
CA	Canada	IT	Italy	NE	Niger	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NL	Netherlands	VN	Viet Nam
CG	Congo	KE	Kenya	NO	Norway	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NZ	New Zealand	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	PL	Poland		
CM	Cameroon	KR	Republic of Korea	PT	Portugal		
CN	China	KZ	Kazakstan	RO	Romania		
CU	Cuba	LC	Saint Lucia	RU	Russian Federation		
CZ	Czech Republic	LI	Liechtenstein	SD	Sudan		
DE	Germany	LK	Sri Lanka	SE	Sweden		
DK	Denmark	LR	Liberia	SG	Singapore		
EE	Estonia						

DYNAMIC O_x BIOLOGICAL BURDEN REDUCTION

FIELD OF THE INVENTION

The present invention generally relates to a method and apparatus that can be utilized to significantly reduce the biological load on consumer products such as food products, botanicals, cosmetic ingredients and medical products, which have traditionally been treated with commercial sterilants or fumigants such as ethylene oxide, propylene oxide, methyl bromide, hydrogen phosphide, steam (heat), irradiation, and the like.

BACKGROUND OF THE INVENTION

Currently, ethylene oxide (EO) is the agent of choice for virtually all of the gaseous sterilization of consumer products performed in the United States. EO, however, has a number of properties which limit its use as a sterilant. For example, EO is highly flammable and highly explosive. For this reason, all EO sterilization facilities must include damage-limiting construction and all equipment used in the EO sterilization process must be explosion proof. In addition, high concentrations of EO are acutely toxic for humans, and EO has been implicated as a carcinogen in laboratory animals. EO sterilization plants must, therefore, be outfitted with expensive equipment to minimize exposure of facility personnel to EO and to prevent escape of EO to the atmosphere outside the facility.

Another disadvantage of EO is that it reacts with chlorine to form ethylene chlorhydrin (ECH) and with water to form ethylene glycol (EG). Both of these byproducts are toxic and must be reduced to safe levels before sterilized materials are released for use. In fact, formation of ECH and EG preclude the use of EO for sterilization of a number of food products for which it would otherwise be a very useful sterilant.

Over the last fifty years, a number of other gases that lack one or more of the disadvantages of EO have been tested as sterilants. These gases include, for example, hydrogen peroxide, ozone and chlorine dioxide. However, no other gas has proven to be as efficacious as EO for use in large volume industrial sterilizers. The most common reason for failure of these gases as useful sterilants has been inadequate permeation of the gases through packaging materials and into lumens and interstices of the product being sterilized. Failure to permeate may be due to a number of factors including size of the sterilant molecule, reactivity of the sterilant molecule with outer packaging materials prior to reaching the target material, and degradation of the sterilant molecule prior to reaching the target material.

A number of commercial fumigants are presently used to treat foodstuffs and other stored commodities. The most widely used fumigants are methyl bromide, hydrogen phosphide, carbon dioxide, and hydrogen cyanide. Many of these compounds pose hazardous conditions for application personnel and can form deleterious residues in the foodstuffs and commodities that are treated. Methyl bromide, the most widely used fumigant, has been identified as an ozone layer depleting agent and is slated, under the Montreal Protocols of 1997, to be banned by the year 2005. One important factor that differentiates the above listed fumigants from other sterilants is their ability to readily permeate granular or powdered products, which allows the fumigant to contact the infesting pests in such products.

Some of the above-mentioned traditional sterilants or fumigants have been identified with the formation of carcinogens and mutagens which thus limit the products that can be treated.

Other procedures that have been developed to treat products utilize heat, ionizing radiation, and other chemical compounds. All of these procedures are potentially detrimental to the products' nutritional, physical and/or chemical attributes and thus make them undesirable.

Thus, a means to reduce biological loads in consumer products to eliminate human pathogens while maintaining product stability remains a necessity. Accordingly, there remains a need for better alternative methods and apparatus for reducing the biological load on consumer products.

SUMMARY OF THE INVENTION

It is desirable to treat a wide variety of consumer products in chamber type operations in a cost effective manner. The method and apparatus of the present invention permit sterilization or fumigation (hereinafter referred to as "biological burden reduction" of a product in its original container (e.g., burlap bag, fiber drum, kraft paper bag, plastic bag, etc.). Thus, double handling, product loss, and post treatment contamination are reduced.

The method of the present invention utilizes a gaseous mixture containing O_1 , O_2 and O_3 (hereinafter referred to as " O_x ") in a technologically advanced treatment system that overcomes the limitations formerly encountered with O_3 treatment on biological burden. Prior O_3 treatments include, for example, (1) the submersion of an article to be sterilized in ozone-containing water and the bubbling of ozonated water over the article (see, e.g., U.S. Patent No. 4,517,159 to Karlson and U.S. Patent No. 4,640,872 to Burleson); and (2) the static treatment of medical devices and food products with gaseous ozone (see, e.g., U.S. Patent No. 3,179,017 to Shapiro et al., U.S. Patent No. 5,069,880 to Karlson, and U.S. Patent

No. 5,120,512 to Masuda.) Systems utilizing ozone as a sterilant such as described above have encountered several limitations. The incorporation of ozone gas into water and then submersion of items(s) to be sterilized or the spraying of ozone treated water onto the surface of item(s) to be sterilized limit the process to products that can be soaked in water. The few gaseous uses of ozone have been limited to the surface treatment of medical devices and the like due to the lack of adequate penetration into compacted products. Thus, although these past processes have proven the efficacy of ozone as a sterilant, the limitation of the use of ozone as a surface treatment has not presented ozone as a reliable sterilant or fumigant for products contained within commercial containers. Furthermore, until recently, high concentration ozone generators, which would have allowed the present invention to properly function, have not been commercially available. The present invention requires a relatively high concentration of O_x for an extended treatment period to assist the required permeation of the O_x into the substrate being treated. In addition to the generation of the ozone molecule, the present invention also utilizes the quenching effect of other inert gases to assist ozone generation, thereby increasing the stability of the O_x radicals. A combination of oxygen, carbon dioxide, argon, and nitrogen have been used in the method of the present invention to achieve these factors. In addition to the benefits discussed above, the use of small quantities of carbon dioxide results in an increase in the rate of respiration in insects and some microbes, thus further aiding the action of the O_x gases. Furthermore, the presence of atmospheric nitrogen has been utilized in the food industry for many years to protect sensitive oils and fats from oxidative rancidity. Small quantities of nitrogen can be used in the method of the present invention to assist in the protection of sensitive food components as well as assisting in the stabilization of the O_x generation.

As an aid to understanding the invention, but without being limited thereby, the present invention is based on the discovery of the following:

1. That water submersion in O₃ sterilization is unnecessary and, in many cases, undesirable;
2. That use of gaseous O₃ in a static fashion, in many cases, does not provide adequate penetration of O₃ into products being treated;
3. That treating a product in-situ (no repackaging needed) is desirable;
4. That the use of cooler temperatures, in many cases, is desirable; and
5. That the destruction of insects and their eggs is desirable.

Accordingly, it is an object of the present invention to provide a method and apparatus for reducing biological burden from consumer products.

It is another object of the present invention to provide a method and apparatus for reducing biological burden from consumer products in a safe manner.

It is thus an object of the present invention to eliminate the health risks that are associated with the reduction of biological burden from consumer products.

It is a further object of the present invention to provide a simple, efficient and economical method and apparatus for reducing biological burden from consumer products that can be used at the site of production and/or packaging of such products.

In accordance with the above and other objects, the inventive method comprises applying a continuous stream of O_x gas to a material in a sealed biological burden reduction chamber. The continuous stream of O_x gas is prepared in an O_x generation cell, which contains a means for generating the O_x gas at a pressure less than about 20 lbs/in² using, for example, one or more of the following: corona discharge, high frequency electrical

discharge, ultraviolet light, x-ray, radioactive isotope and electron beam.

As discussed herein, a small percentage of N₂, CO₂ and/or Ar may be added during O_x treatment. The addition of 0% to 15% N₂, 1% to 20% CO₂ and/or 1% to 18% Ar increases the generation of an O_x quenching effect. Penetration of O_x into the material being treated is thus enhanced. In addition, Argon is unique among the (inert) Noble Gases, in that it is soluble in both water and organic liquids. (The Merck Index Eleventh Edition). This characteristic theoretically enables Argon to become a glue of sorts. Argon is capable of attaching to gases without reacting thereto. Argon thus assists in O_x quenching by attaching to the O_x molecules and preventing the O_x molecules from colliding into each other. Argon also loosely binds hydrophilic and hydrophobic materials, thus allowing one to be diffused through the other, without reacting with either. This characteristic is useful in accelerating the diffusion of O_x into and through hydrophilic materials such as fats, oils and cell walls.

The inventive apparatus comprises:

- (a) a biological burden reduction chamber;
- (b) a vacuum pump coupled to the biological burden reduction chamber;
- (c) an O_x generation cell, wherein the O_x generation cell contains a means for generating O_x at pressure less than about 20 lbs./in² using, for example, one or more of the following: corona discharge, high frequency electrical discharge, ultraviolet light, x-ray, radioactive isotope and electron beam;
- (d) a first control valve coupled to the biological burden reduction chamber and the O_x generation cell, wherein the first control valve is capable of permitting O_x to be drawn from the O_x generation cell into the biological burden reduction chamber; and
- (e) a second control valve coupled to the biological burden reduction

chamber, wherein the second control valve is capable of withdrawing O_x contained within the biological burden reduction chamber out of the biological burden reduction chamber.

Water vapor may be introduced to the gaseous O_x to maintain an appropriate humidity level, i.e., between about 20% and 98% relative humidity, and, more preferably between about 40% and 75% relative humidity. The appropriate humidity level is dependent upon the ambient humidity and upon the product being treated. For example, granular and powdered products require a relatively low humidity level to prevent growth of mold and yeast thereon. However, depending on the length of treatment time, the vacuum created during the process removes humidity, thus requiring the addition of humidity. The O_x gas may then be passed through a commercially available catalytic destruct unit to eliminate any residual O_3 and O_1 before the gas stream is discharged to the atmosphere.

The present invention is also directed to treated consumer products that result from use of the present inventive method and apparatus.

Additional objects and attendant advantages of the present invention will be set forth in the description and examples that follow, or may be learned from practicing the method or using the apparatus of the present invention. These and other objects and advantages may be realized and attained by means of the features, instrumentalities and/or combinations particularly described herein. It is also to be understood that the foregoing general description and the following detailed description are only exemplary and explanatory and are not to be viewed as limiting or restricting the invention as claimed.

The invention itself, together with further objects and attendant advantages, will best be understood by reference to the following detailed description, taken in conjunction with the accompanying drawings.

BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1 is a flow chart showing an embodiment of a method for using a continuous flow of O_x to reduce biological burden in accordance with the method of the present invention.

FIG. 2 is a schematic showing one example of an apparatus for using a continuous flow of O_x to reduce biological burden in accordance with the method of the embodiment in FIG.1.

FIG. 3 is a graph comparing permeation of O_x gas for (a) VVP + CO_2 + O_x in accordance with the method of the invention, (b) VVP + O_x in accordance with the method the invention, and (c) static + O_x in accordance with conventional use of gaseous O_x in a static fashion .

In the following description, like parts are designated by like reference numerals throughout the figures.

DESCRIPTION OF PREFERRED EMBODIMENTS

All patents, patent applications and literatures that may be cited herein are incorporated herein by reference.

The antibacterial potential of O_3 has been recognized for many years. O_3 is widely used as a disinfectant for sewage treatment and for purification of drinking water. It has, however, failed to gain acceptance as a biological burden reduction treatment for medical supplies and consumer goods. The primary reason for this failure is that the O_3 molecule is highly unstable and quickly reverts to O_2 if it does not encounter a susceptible substrate with which to react. Ozone also has the capacity to react with a broad array of substrates and

would be expected to react with packaging materials surrounding the items being sterilized. This further reduces the number of O_3 molecules available to react with and inactivate microbial contaminants.

Previous attempts to use O_3 as a biological burden reduction treatment include the reliance upon filling a sterilization chamber with O_3 and exposing the materials to be treated in static fashion for various periods of time without replenishment of O_3 . See for example, U.S. Patent Nos. 3,719,017 and 5,069,880. Under these conditions, the concentration of O_3 within the chamber would be expected to rapidly decrease to a level below that required for effective biological burden reduction due to the short half life of O_3 , which is typically less than 20 minutes. A further disadvantage of the static exposure technology is the reliance on simple diffusion to promote permeation of the O_3 molecules through packaging materials and into voids and interstices of the materials being treated. Thus, such methods do not achieve adequate permeation of into the material being treated.

The present invention, which has been designated "dynamic O_x biological burden reduction," offers significant advances over the prior static biological burden reduction technology in that it provides a uniformly high concentration of O_x , i.e., between about 3% and 16%, throughout the treatment cycle and promotes rapid permeation of O_x through packaging materials and into the voids and interstices of the materials undergoing treatment. Continuous operation of the vacuum pump and O_x generator during biological burden reduction ensures that the concentration of O_x remains high throughout the process by constantly supplying newly generated O_x molecules to replace those molecules which have spontaneously degraded to inactive O_2 and those which have reacted during the process. Operation of the vacuum pump during the process, as practiced in the present invention, also

maintains a continuous pressure differential, which actively draws O_x molecules through packaging materials and into the materials being treated.

Dynamic O_x biological burden reduction provides significant cost advantages over existing biological burden reduction technology. The most significant savings derive from the fact that the O_x biological burden reducing gas may be generated on site, during the process.

Because O_x is not flammable or explosive, facilities need not include damage-limiting construction or explosion-proof equipment. Another advantage of dynamic O_x biological burden reduction is that scrubbing will be easily accomplished using existing technology. Moreover, O_3 is classified by the U.S. Food and Drug Administration as a generally recognized as safe "GRAS" substance.

The dynamic O_x biological burden reduction process of the invention has proven successful in the treatment of a wide variety of materials, including medical products, spices, flavorings, and packaging materials.

Referring to FIG. 3, which is a graph comparing permeation of O_x gas for (a) VVP + CO_2 + O_x in accordance with the method of the invention, (b) VVP + O_x in accordance with the method the invention and (c) static + O_x in accordance with conventional use of gaseous O_x in a static fashion. The static flow of gas (c) demonstrated a limited permeation, which quickly stopped altogether. The (VVP) process (a) and (b), on the other hand, demonstrated a continuous progression through the packed column; completely depleting the chemical indicator. The method of the present invention enhances the permeability of O_x gases into commercially sized containers of granular and powdered food components. The observation of live insects and microbiological and chemical indicators have been utilized to measure the

incorporation of lethal doses of O_x into these containers. Via comparative data (see FIG. 3), a static flow of O_x bearing gas (curve (c)) has shown ineffective in driving the O_x into the containers. The method of the present invention utilizes a process herein described as the **Vacuum Vapor Phase Dynamic Flow (VVP)**. In theory, and supported by empirical evidence, VVP acts as the driving force to enhance permeation of the O_x gases by two factors. The first factor is the molecular acceleration of the O_x gases due to the flashing of the concentrated O_x gas into the reduced pressure treatment chamber. This creates a driving force at a molecular level that continuously forces the O_x gases into the product being treated. The second factor is the resulting reduction of molecules within the reduced pressure treatment chamber which reduces the incidence of molecular collision of the O_x gases. Molecular collision of the O_x gases causes rapid degradation of the O_3 and O_1 radicals present therein, thereby reducing the gases' effectiveness. Without the VVP process, the O_x gas flow could only be utilized as a surface treatment of non-amorphous materials, therefore, the VVP process expands the capabilities of the present invention to process virtually any type of product in-situ, thus eliminating the need to repackage the product after treatment.

The method of the present invention avoids many of the limitations of previous practices by avoiding the need for water sprays and/or water immersion of the substrate to be treated. Many products such as spices, flour-based products, sugar-based products, cosmetic bases, herbs, and botanicals, which are sensitive to high levels of moisture, can be treated using the method of the present invention. The method of the present invention also avoids the need to open conventional commercial packaging before treatment, thus avoiding unnecessary product degradation and loss. The product is treated in situ utilizing a

conventional metal chamber with the product palletized in its normal shipping configuration. Previous methods have required the product to be agitated, blended, bubbled, or re-packaged during or immediately upon completion of the treatment. Due to the increased permeation of the VVP process and the O_x gas mixture, these damaging handling practices are avoided. The unique gas mixture also assists in the stabilization of the O_x generation through its quenching effect. The extended half life of the O_x radicals allows the active portions of the treatment gas to fully penetrate the substrate and act upon offending organisms. In combination with carbon dioxide, the stabilized O_x gas mixture is further enhanced by the increased respiration rates of the offending organism(s) while in the presence of the permeated O_x gases. Traditional fumigants and sterilants have required elevated temperatures for prolonged periods of time to be effective. In the case of heat sensitive products such as fruits and spices, the shelf life and/or quality of the product can be severely compromised during such treatment. The method of the present invention protects such products by processing them under relatively cool conditions, i.e, between about 32°F and 80°F.

Referring to FIG. 2, according to an embodiment of the invention, an apparatus according to the invention may include a biological burden reduction chamber 1 equipped with a gasketed door 2 that can be opened to accommodate placement of material 3 within the biological burden reduction chamber 1 and tightly closed and latched. The biological burden reduction chamber 1 permits a vacuum tight seal during the process. The chamber 1 is connected via piping and appropriate control valves 4 to a vacuum pump 5 and separately to a generator of O_x 6, which, in turn is connected to a gas washer 14 and an air preparation regulated feed gas supply 15. The biological burden reduction chamber 1 is jacketed by

coils of metal tubing 11 through which heated or chilled water generated by a temperature control (e.g., glycol) system 7 may be pumped to regulate the temperature within the chamber 1 during the biological burden reduction process. The entire biological burden reduction process may be controlled and monitored by a programmable industrial process controller 8. The chamber 1 is also connected to a water vapor source 12 to provide humidity control.

According to the method of the embodiment, material 3 for which biological burden is to be reduced is placed within the biological burden reduction chamber 1 and the door 2 is closed and latched. The process is then initiated by activating the process controller 8, which has previously been programmed with the appropriate process parameters such as pressure, temperature and humidity. The controller 8 first activates the vacuum pump 5 and ancillary valves 4 to reduce the biological burden reduction chamber pressure to a preset level between, e.g., 0 and 20 lbs/in² depending on the pressure sensitivity of the product being treated, to introduce via the water vapor source 12 the desired humidity, and to maintain a desired temperature via the temperature control system 7. After the appropriate vacuum level has been reached, the controller 8 initiates biological burden reduction by activating the O_x generator 6 and opening a control valve 10, allowing the washed O_x stream to be drawn into, through and out of the chamber 1 by the pressure differential. The vacuum pump 5 and O_x generator 6 operate continuously during the process.

Exposure to the O_x gas mixture may be varied in time from several minutes to several hours, depending on the material being treated. Once the biological burden reduction phase is complete, the vacuum pump 5 and O_x generator 6 are inactivated and fresh air is allowed to enter the chamber 1 via the air purge valve 13. All O_x gases may then be passed through a

commercially available catalytic destruct unit 9 which eliminates any residual O_3 and O_1 before the gas stream is discharged to the atmosphere. The treated material 3 can then be removed from the chamber 1 and is ready for use following appropriate tests to confirm biological burden reduction.

The present invention will be further illustrated by the following non-limiting Examples.

Example 1

O_3 Treatment of Black Pepper (*Piper nigrum* L.) In situ (50 kg burlap bags)

(1) Parameters

O_3 Concentration (by weight)	: 5.4%
CO_2 Concentration (by weight)	: 6.0%
Argon Concentration (by weight)	: 2.5%
Chamber Temperature	: 12° C
Relative Humidity	: 64%
Time (VVP)	: 16 hours
Pressure	: 5.8 p.s.i.a.

(2) Results

	<u>Control</u>	<u>VVP Treated</u>
Total Plate Count	5,500,000	100,000
Total Coliforms	5,000	<10
Escherichia coli	600	<10
Yeast spp.	1,500	<10
Mold spp.	4,500	<10
Bacillus subtilis (spores)	1,600,000	<10

Example 2**O₃ Treatment of Echinacea purpurea In situ (50 kg plastic woven bags)****(1) Parameters**

O ₃ Concentration (by weight)	: 4.8%
CO ₂ Concentration (by weight)	: 9.5%
Argon Concentration (by weight)	: 3.0%
Chamber Temperature	: 17° C
Relative Humidity	: 70%
Time (VVP)	: 20 hours
Pressure	: 4.7 p.s.i.a.

(2) Results

	<u>Control</u>	<u>VVP Treated</u>
Total Plate Count	2,300,000	180,000
Total Coliforms	12,000	100
Escherichia coli	1,200	<10
Yeast spp.	3,400	<10
Mold spp.	2,600	<10
Bacillus subtilis (spores)	1,600,000	<10

Example 3**O₃ Treatment of Milk Thistle Seed In situ (80 pound plastic woven bags)****(1) Parameters**

O ₃ Concentration (by weight)	: 4.5%
CO ₂ Concentration (by weight)	: 5.0%
Argon Concentration (by weight)	: 1.0%
Chamber Temperature	: 20° C
Relative Humidity	: 60%
Time (VVP)	: 24 hours
Pressure	: 3.5 p.s.i.a.

(2) Results

	<u>Control</u>	<u>VVP Treated</u>
Total Plate Count	4,700,000	210,000
Total Coliforms	32,000	<10
Escherichia coli	5,800	<10
Yeast spp.	4,500	<10
Mold spp.	16,000	<10
Bacillus subtilis (spores)	1,600,000	<10

Example 4**O₂ Treatment of Alfalfa Seed In situ (50 pound canvas bags)****(1) Parameters**

O₂ Concentration (by weight) : 3.5%
CO₂ Concentration (by weight) : 1.0%
Argon Concentration (by weight) : 1.0%
Chamber Temperature : 22° C
Relative Humidity : 30%
Time (VVP) : 12 hours
Pressure : 7.5 p.s.i.a.

(2) Results

	<u>Control</u>	<u>VVP Treated</u>
Total Plate Count	3,800,000	220,000
Total Coliforms	18,000	<10
E. coli 0157:H	75,200	<3
Bacillus subtilis (spores)	1,600,000	<10
Germination (%)	96	92

Example 5**O₂ Treatment of Navel Oranges for Bean Thrips, Red Scale, and Fuller Rose Beetle.****(1) Parameters**

O₂ Concentration (by weight) : 4.0%
CO₂ Concentration (by weight) : 10.0%
Argon Concentration (by weight) : 1.0%
Chamber Temperature : 20° C
Relative Humidity : 30%
Time (VVP) : 2 hours
Pressure : 9.5 p.s.i.a.

(2) Results

All adults larvae, and eggs (Fuller Rose Beetle only) were killed in the process. The treated fruit was incubated for 28 days after treatment to assure that all three life cycles had been destroyed.

The *Bacillus subtilis* spore strips are standardized control tabs that are produced by Amsco Medical Products, Erie, Pennsylvania, and are used as standards for commercial sterilization processes.

EO and heat have been utilized in the past to de-germ various feed seeds to eliminate the possibility of germination where it is undesirable or would pose a weed growth problem. Certain seed (e.g., safflower), which have been found to be sensitive to EO gas are readily de-germed utilizing the present invention. By controlling certain parameters, some seed types (e.g., alfalfa) have been found to be very resistant to damage by O_x gases, which would lend the present invention to process these seeds for microbiological reduction while still maintaining a high percentage of germination.

The dynamic biological burden reduction process of the invention is thus an excellent substitute for commercial sterilants and fumigants in all of its current uses and is also useful for the treatment of many food ingredients on which commercial sterilant and fumigant use are not permitted, including cocoa beans, grains, and edible gums.

The dynamic O_x biological burden reduction process has also been shown to be highly insecticidal and is therefore a useful substitute for certain current uses of methyl bromide, which, as discussed herein, is soon to be banned under the direction of the Montreal Protocols of 1997.

WHAT IS CLAIMED:

- 1 1. A method for biological burden reduction, comprising a step of applying a
2 continuous stream of O_x to a material in a sealed biological burden reduction chamber,
3 wherein said O_x includes oxygen and its radicals.
- 1 2. The method of claim 1, wherein O_x is selected from an integer from 1 to 3.
- 1 3. The method of claim 1, further comprising continuously withdrawing O_x
2 from said sealed biological burden reduction chamber.
- 1 4. The method of claim 1, further comprising creating a pressure differential
2 within said biological burden reduction chamber and maintaining said pressure differential
3 while continuously applying said stream of O_x to said material.
- 1 5. The method of claim 4, further comprising agitating said O_x in said biological
2 burden reduction chamber to increase permeation of said O_x into said material.
- 1 6. The method of claim 5, wherein forced air is used to agitate said O_x .
- 1 7. The method of claim 5, wherein said agitating distributes said O_x evenly
2 throughout said biological burden reduction chamber.
- 1 8. The method of claim 5, further comprising
 - 2 (a) creating a vacuum within said biological burden reduction chamber;
 - 3 (b) generating O_x in an O_x generation cell;
 - 4 (c) withdrawing a stream of O_x from said O_x generation cell into said
5 biological burden reduction chamber; and
 - 6 (d) withdrawing O_x from said biological burden reduction chamber.
- 1 9. The method of claim 4, wherein said O_x generation cell comprises an O_x
2 generator capable of generating O_x at a pressure of less than 20 lbs/in² selected from one or

1 more of the group consisting of corona discharge, high frequency electrical discharge,
2 ultraviolet light, x-ray, radioactive isotopes and electron beam.

1 10. The method of claim 8, wherein said O_x in said biological burden reduction
2 chamber is maintained at a concentration of about 0.1% to about 100% per total volume of
3 gases in said biological burden reduction chamber.

1 11. The method of claim 10, wherein O_3 in said biological burden reduction
2 chamber is maintained at a concentration of about 0.1% to about 25% per total weight of
3 gases in said biological burden reduction chamber.

1 12. The method of claim 11, wherein said O_3 in said biological burden reduction
2 chamber is maintained at a concentration of about 3% to about 16% per total weight of gases
3 in said biological burden reduction chamber, wherein an amount of O_3 used is dependent on
4 said material.

1 13. The method of claim 8, further comprising maintaining a pressure differential
2 between a pressure within said O_x generation chamber and a pressure within said biological
3 burden reduction chamber sufficient to continuously withdraw said O_x through said
4 biological burden reduction chamber.

1 14. The method of claim 8, further comprising using a biological burden
2 reduction chamber of about 100 ft.³ to about 8000 ft.³.

1 15. The method of claim 8, further comprising using a biological burden
2 reduction chamber of about 1 ft.³ to about 100 ft.³.

1 16. The method of claim 13, further comprising controlling water vapor present
2 in said continuous stream of O_x prior to applying said continuous stream of O_x to said
3 material.

- 1 17. The method of claim 12, wherein said pressure within said biological burden
2 reduction chamber is maintained between about 0 psia and 20 psia.
- 1 18. The method of claim 1, wherein said O_x is generated from ambient air or
2 components of ambient air.
- 1 19. The method of claim 1, wherein said O_x is generated from other oxygen
2 sources including gaseous oxygen, liquid oxygen, H_2O and mercuric oxide.
- 1 20. The method of claim 1, wherein the material is a food product.
- 1 21. The method of claim 1, wherein the material is a medical product.
- 1 22. The method of claim 1, wherein the material is a cosmetic ingredient.
- 1 23. The method of claim 1, wherein the material is a dietary supplement.
- 1 24. The method of claim 1, wherein the material is a botanical.
- 1 25. The method of claim 1, wherein the material is a nutraceutical.
- 1 26. The method of claim 1, wherein the material is a pharmaceutical ingredient.
- 1 27. The method of claim 1, wherein the material is a packaging material.
- 1 28. The method of claim 1, wherein the material is a nursery stock product.
- 1 29. The method of claim 1, wherein the material is a color additive.
- 1 30. The method of claim 1, wherein the material is a seed.
- 1 31. The method of claim 1, wherein the material is a personal care product.
- 1 32. The method of claim 1, wherein the material is an animal feed.
- 1 33. The method of claim 1, wherein the material is a flavoring.
- 1 34. An apparatus for biological burden reduction, comprising:
2 (a) a biological burden reduction chamber;
3 (b) a vacuum pump coupled to said biological burden reduction chamber;

4 (c) an O_x generation cell, wherein said O_x generation cell comprises a
5 means for generating O_x ;

6 (d) a first control valve coupled to said biological burden reduction
7 chamber and said O_x generation cell, wherein said first control valve is capable of permitting
8 said O_x to be withdrawn from said O_x generation cell into said biological burden reduction
9 chamber; and

10 (e) a second control valve coupled to said biological burden reduction
11 chamber, wherein said second control valve is capable of withdrawing O_x contained within
12 said biological burden reduction chamber.

1 35. The apparatus of claim 34, further comprising a member for creating forced
2 air contained within said biological burden reduction chamber, wherein said forced air
3 distributes said O_x evenly throughout said biological burden reduction chamber.

1 36. The apparatus of claim 34, further comprising a temperature-regulating
2 means.

1 37. The apparatus of claim 34, further comprising a means for controlling water
2 vapor coupled to said biological burden reduction chamber.

1 38. The apparatus of claim 34, further comprising a controller for controlling and
2 monitoring physical parameters within said biological burden reduction chamber.

1 39. The method of claim 1, wherein said biological burden is selected from a
2 group of living entities including insects, bacteria, viruses, algae, yeasts, molds, nematodes,
3 parasites and weed seed.

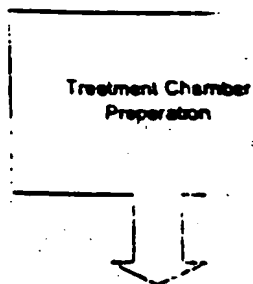
1 40. The apparatus of claim 36, further comprising a means to convert said O_x to
2 O_2 prior to release into atmosphere.

1 41. The method of claim 17, wherein a humidity of an atmosphere within said
2 biological burden reduction chamber is between about 20% to about 98%.

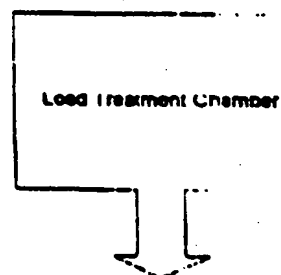
1 42. The method of claim 4, wherein a temperature within said biological burden
2 reduction chamber is between about 32°F and about 80°F.

1 43. The method of claim 1, wherein a flow rate of said continuous stream of O_x
2 within said biological burden reduction chamber is between about 0.1 L/min/ft³ and about 2
3 L/min/ft³.

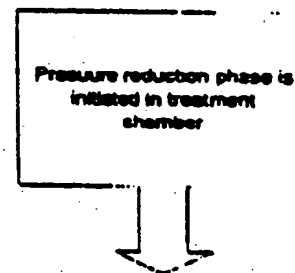
1 44. The method of claim 1, further comprising applying a continuous stream of
2 one or more of a gas selected from the group consisting of N₂, CO₂ and Ar in addition to
3 said continuous stream of O_x.



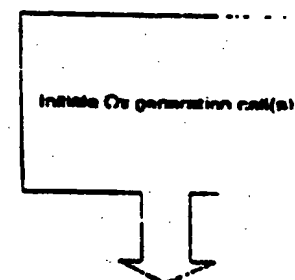
The treatment chamber temperature, pressure, and relative humidity are stabilized to the ideal treatment parameters for the product to be treated.



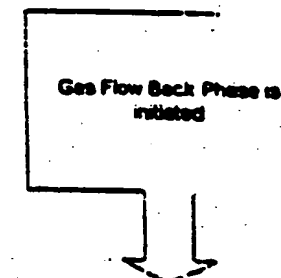
The product to be treated is placed into the treatment chamber. The door with an airtight seal is closed and the seal is engaged. The product is allowed to equilibrate before processing.



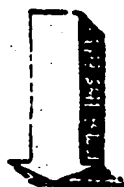
An initial pressure reduction is initiated within the treatment chamber to remove competing gases.



A properly dried and mixed gas mixture is prepared to feed through the O₂ generation cell(s) to generate the O₂ gas flow.

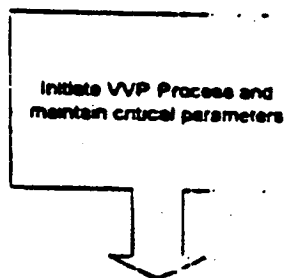


The ancillary control valve is adjusted to allow a controlled flow of the O₂ gas into the treatment chamber. The pressure, temperature, O₂ concentration, and water vapor concentration are controlled.

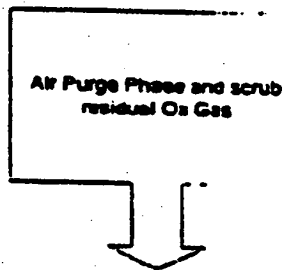




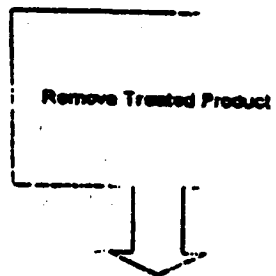
Continued from Page 1



The continuous flow VVP phase is allowed to continue for a predetermined amount of time to assure permeation into the product being treated. The pressure, temperature, Ox concentration, and water vapor are carefully monitored and controlled.



The Ox flow is terminated and the treatment chamber pressure is cycled to flush the treatment chamber of residual Ox gas.



The treated product is removed from the treatment chamber ready for further processing or shipment.

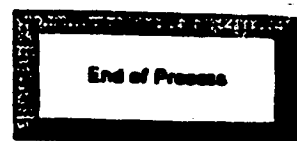


FIGURE 1

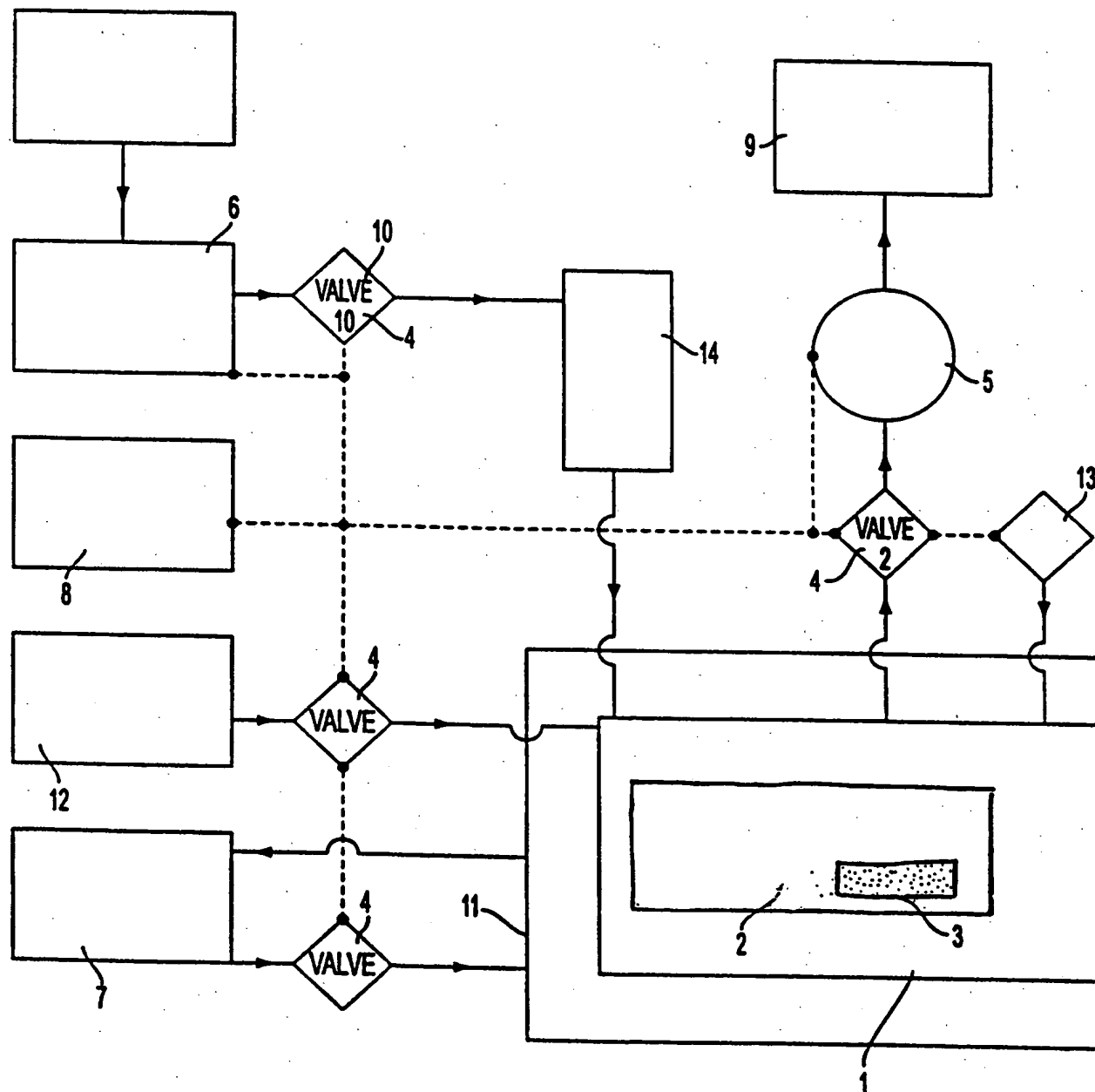


FIG. 2

SUBSTITUTE SHEET (RULE 26)

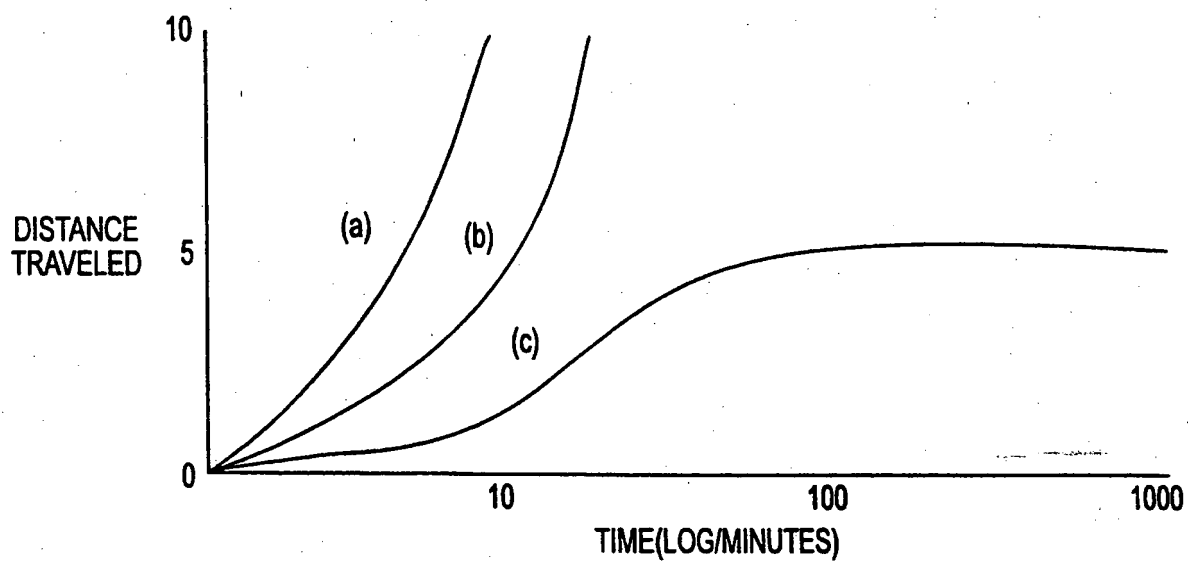


FIG. 3

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US98/26232**A. CLASSIFICATION OF SUBJECT MATTER**

IPC(6) : A61L 2/20; B01J 19/08.

US CL : 422/186.12, 29, 33.

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 422/186.12, 29, 33.

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)
APS.**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 5,069,880 A (KARLSON et al.) 03 December 1991, see abstract and Fig.8.	34-40
A	US 5,120,512 A (MASUDA) 09 June 1992, see abstract.	1-33 and 41-44

☐ Further documents are listed in the continuation of Box C. ☐ See patent family annex.

* Special categories of cited documents:	*T	later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
A document defining the general state of the art which is not considered to be of particular relevance	*X*	document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
B earlier document published on or after the international filing date	*Y*	document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
L document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	*A*	document member of the same patent family
O document referring to an oral disclosure, use, exhibition or other means		
P document published prior to the international filing date but later than the priority date claimed		

Date of the actual completion of the international search

11 MARCH 1999

Date of mailing of the international search report

01 APR 1999

Name and mailing address of the ISA/US
Commissioner of Patents and Trademarks
Box PCT
Washington, D.C. 20231

Facsimile No. (703) 305-3230

Authorized officer

KISHOR MAYEKAR

Telephone No. (703) 308-0661

Form PCT/ISA/210 (second sheet)(July 1992)*

THIS PAGE BLANK (USPTO)